

1. A polypeptide construct comprising:
 - a first portion comprising the second constant domain (C_{H2}) and/or third constant domain (C_{H3}) of an antibody heavy chain, and
 - a second portion comprising at least two TGF- β superfamily receptor ectodomains (T β SR-ED) linked in tandem,
 wherein the N-terminus of the second portion is linked to the C-terminus of the first portion.
2. A polypeptide construct comprising:
 - a first portion comprising the second constant domain ($CH2$) and/or third constant domain ($CH3$) of an antibody heavy chain, and
 - a second portion comprising at least one TGF- β superfamily receptor ectodomains (T β SR-ED),
 wherein the N-terminus of the second portion is linked to the C-terminus of the first portion, and further wherein the first portion does not further comprise an antibody that binds to an antigen that is PD-L1, EGFR1, Her-2, CD4, CD6, CD20, CD25, MUC-1, IL-2, IL-6, or CTLA-4.
3. A polypeptide construct comprising:
 - a first portion comprising the second constant domain (C_{H2}) and/or third constant domain (C_{H3}) of an antibody heavy chain, and
 - a second portion comprising at least one TGF- β superfamily receptor ectodomain (T β SR-ED),
 wherein the N-terminus of the second portion is directly fused to the C-terminus of the first portion.
4. A polypeptide construct comprising
 - a first portion comprising the second constant domain (C_{H2}) and/or third constant domain (C_{H3}) of an antibody heavy chain, and
 - a second portion comprising at least one TGF- β superfamily receptor ectodomain (T β SR-ED),
 wherein the N-terminus of the second portion is linked to the C-terminus of the first portion, and wherein the polypeptide construct neutralizes TGF- β with at least 100-fold more potency than the T β SR-ED alone.
5. The polypeptide construct of claims 2-4, wherein the second portion comprises one T β SR-ED.
6. The polypeptide construct of claim 5, wherein the second portion comprises two T β SR-EDs.
7. The polypeptide construct according to claims 1-6, wherein the T β SR-ED is a TGF- β receptor type II ectodomain (T β R-II-ED).
8. The polypeptide construct of claims 1-6, wherein the T β SR-ED comprises a sequence selected from the group consisting of SEQ ID NO:35, SEQ ID NO:69, SEQ ID NO:75, SEQ ID NO:81, and a sequence substantially identical thereto.
9. The polypeptide construct of claims 1-8, wherein the second portion comprises a sequence selected from the group consisting of SEQ ID NO:43-SEQ ID NO:51, SEQ ID NO:61-SEQ ID NO:68, SEQ ID NO:73, SEQ ID NO:74, SEQ ID NO:79, SEQ ID NO:80, SEQ ID NO:85, SEQ ID NO:86, SEQ ID NO:88, and a sequence substantially identical thereto.
10. The polypeptide construct of any one of claims 1-8, wherein the first portion further comprises a C_{H1} , a C_{H1} and V_{H^*} or C_{H1} and scFv.
11. The polypeptide construct of any one of claims 1-10, wherein the antibody heavy chain is of human origin.
12. The polypeptide construct of any one of claims 1-11, wherein the antibody heavy chain is selected from the group consisting of a human IgG1, IgG2, IgG3, or IgG4 heavy chain.
13. The polypeptide construct of any one of claims 1-12, wherein the antibody heavy chain is a human IgG1.
14. The polypeptide construct of claim 4, wherein the polypeptide construct shows longer in vivo half-life compared to the half-life of the second portion alone.
15. The polypeptide construct of any one of claims 1-14, wherein the polypeptide construct is a single chain polypeptide.
16. The polypeptide construct of any one of claims 1-15, wherein the polypeptide construct forms a dimeric polypeptide.
17. The polypeptide construct of claims 1-16, wherein the polypeptide construct is heterodimeric.
18. A polypeptide construct selected from the group consisting of any one of SEQ ID NO:91 to SEQ ID NO:120, and a sequence substantially identical thereto.
19. A polypeptide construct according to claims 1-16, wherein the construct comprises an antibody, antigen binding fragment thereof, or a targeting moiety.
20. A polypeptide construct according to claim 19, comprising the antibody, antigen binding fragment, or targeting moiety at the N-terminus of the first portion.
21. A polypeptide construct according to claim 19, wherein the antigen binding fragment may be selected from the group consisting of a Fv, scFv, Fab, or sdAb.
22. A polypeptide construct according to claim 19, wherein the antigen binding fragment binds to an antigen that is not PD-L1, EGFR1, Her-2, CD4, CD6, CD20, CD25, MUC-1, IL-2, IL-6, or CTLA-4.
23. A polypeptide construct according to claim 19, wherein the antibody is selected from the group consisting of Cetuximab, Avastin, Herceptin, Synagis, and FC5.
24. A polypeptide construct according to claim 23, wherein the antibody is Cetuximab.
25. A polypeptide construct according to claim 19, wherein the targeting moiety comprises a poly-aspartate sequence motif for bone targeting.
26. A polypeptide construct according to claim 25, wherein the targeting moiety comprises D10.
27. A polypeptide construct according to any preceding claim wherein the construct is a dimeric polypeptide.
28. A polypeptide construct according to claim 27, wherein the dimeric polypeptide comprises:
 - a first single chain polypeptide comprising a first portion comprising the second constant domain (C_{H2}) and third constant domain (C_{H3}) of an antibody heavy chain, and a heavy chain variable region of a given antibody;
 - a second portion comprising one or more TGF- β superfamily receptor ectodomains (T β SR-ED),
 wherein the N-terminus of the second portion is linked to the C-terminus of the first portion, and
 - a second single chain polypeptide comprising a first portion comprising the second constant domain (C_{H2}) and third constant domain (C_{H3}) of an antibody heavy chain, and a light chain variable region of said given antibody;
 - a second portion comprising one or more TGF- β superfamily receptor ectodomain (T β SR-ED) which is the same or different from the ectodomain(s) in the first